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CONCERNING A FILING UNDER 35 U.S.C. 371

U.S. APPLICATION NO. (if known, see 37 CFR 1.5)

09/914066

INTERNATIONAL APPLICATION NO.
PCT/JP00/00862INTERNATIONAL FILING DATE
16 February 2000PRIORITY CLAIMED
23 February 1999

TITLE OF INVENTION

SEAM SOFT CAPSULE PREPARATIONS CONTAINING DIHYDROBENZOFURAN DERIVATIVE

APPLICANT(S) FOR DO/EO/US

Hisakazu KATSUKI

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ The US has been elected in a Demand by the expiration of 19 months from the priority date (PCT Article 31).
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ is attached hereto (required only if not transmitted by the International Bureau).
 - b. ☒ has been communicated by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been communicated by the International Bureau.
 - c. ☐ have not been made, however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
8. ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3))
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ An English language translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An Assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A FIRST preliminary amendment.
 - ☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:
 - ☒ Courtesy copy of the first page of the International Publication (WO 00/50029).
 - ☒ Courtesy copy of the Exact Translation of the International Preliminary Examination Report. There were no annexes.
 - ☒ Formal drawings, 1 sheet, Figures 1-2.
 - ☒ Courtesy Copy of the International Search Report.
 - ☒ The application is (or will be) assigned to: CHUGAI SEIYAKU KABUSHIKI KAISHA, whose address is 5-1, Ukima 5-chome, Kita-ku, Tokyo 115-8543 Japan.

Form PTO-1390 (as slightly revised by Browdy and Neimark)

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:)	Art Unit:
Hisakazu KATSUKI et al.)	
)	
IA No.: PCT/JP00/00862)	
)	Washington, D.C.
IA Filed: February 16, 2000)	
)	
U.S. App. No.:)	
(Not Yet Assigned))	
)	August 23, 2001
National Filing Date:)	
(Not Yet Received))	
)	
For: SEAM SOFT CAPSULE...)	Docket No.: KATSUKI=1

PRELIMINARY AMENDMENT

Honorable Commissioner for Patents and Trademarks
Washington, D.C. 20231

Sir:

Contemporaneous with the filing of this case and
prior to calculation of the filing fee, kindly amend as
follows:

IN THE SPECIFICATION

After the title please insert the following
paragraph:

REFERENCE TO RELATED APPLICATIONS

--The present application is the national stage
under 35 U.S.C. §371 of international application
PCT/JP00/00862, filed February 16, 2000 which designated the
United States, and which application was not published in the
English language.--

REMARKS

The above amendment to the specification is being made to insert reference to the PCT application of which the present case is a U.S. national stage.

Favorable consideration and allowance are earnestly solicited.

Respectfully submitted,
BROWDY AND NEIMARK, P.L.L.C.
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SPECIFICATION

SEAMED SOFT CAPSULE FORMULATION CONTAINING
DIHYDROBENZOFURAN DERIVATIVETECHNICAL FIELD

- 5 The present invention relates to a formulation containing a 5-acetoxy or hydroxy-4,6-di-tert-butyl-2,3-dihydrobenzofuran derivative, especially BO-653, and having an excellent stability and a capability of being absorbed by body.

10

BACKGROUND ART

- A family of 5-acetoxy or hydroxy-4,6-di-tert-butyl-2,3-dihydrobenzofuran derivatives including BO-653, i.e., 4,6-di-tert-butyl-2,2-di-n-pentyl-5-hydroxy-2,3-
- 15 dihydrobenzofuran, are antioxidative compounds, which are generally oily and high-viscous, and thus promise to be useful for treating an arteriosclerosis and the like (WO94/08930). Generally, if such a compound is used as a drug substance for preparing a formulation, it will be
- 20 required to confer a high stability, a capability of being absorbed by body and the like to the compound.

- For example, as a method for preventing the oxidation of a compound such as vitamin E, which is known as a natural antioxidant, it has been reported that the compound
- 25 is combined with another antioxidant or dissolved in a fat and oil such as unsaturated higher fatty acid having double bond(s) and then filled into a soft capsule (JP 7-58082A/95 and JP 8-175967A/96, respectively).

On the other hand, as a method for improving the capability of being absorbed by body of a drug which is highly fat-soluble and slightly water-soluble, it has been reported that the drug is combined with a solubilizer such as a non-ionic surfactant to enhance its water-solubility (JP 7-242535A/95) or is dissolved in a fat and oil such as a medium chain fatty acid triglyceride (JP 8-92088A/96) or a higher unsaturated fatty acid (JP 7-258082A/95).

However, since it is common that a structural difference between compounds is responsible for any differences between them in chemical, physical and/or biological properties, these methods would not be applied to the above-mentioned dihydrobenzofuran derivative, especially BO-653.

Therefore, it would be desirable to provide a formulation which specifically confers a high stability and a capability of being absorbed by body to the derivative.

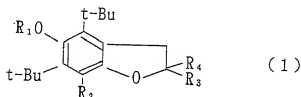
DISCLOSURE OF THE INVENTION

An object of the present invention is to provide a formulation containing a 5-acetoxy or hydroxy-4,6-di-tert-butyl-2,3-dihydrobenzofuran derivative, especially BO-653, or probucol which formulation confers a high stability and a capability of being absorbed by body to the derivative or probucol.

As a result of extensive research, we found that, when the dihydrobenzofuran derivative or probucol is dissolved in a fat and oil without any additional

antioxidant and then filled into a seamed soft capsule, not only its stability is maintained but also its capability of being absorbed by body is improved.

- Accordingly, the present invention provides a seamed soft capsule formulation in which a solution of dihydrobenzofuran derivative represented by general formula (1):



- wherein R₁ represents a hydrogen atom or an acyl group; R₂ represents a hydrogen atom or a lower alkyl group; and R₃ and R₄, which may be identical or different, represent a hydrogen atom or a substituted or unsubstituted alkyl, alkenyl, alkynyl or aryl group, or probucol in a fat and oil, is filled into a seamed soft capsule.

- The present invention also provides a seamed soft capsule formulation containing BO-653. More specifically, the present invention provides a seamed soft capsule formulation in which a solution of BO-653 in fat and oil is filled into a seamed soft capsule.

20 BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 shows the influence of oxygen on the stability of BO-653 in various fat and oils.

Fig. 2 shows plasma BO-653 levels after a solution of BO-653 in various solvents is orally administered.

PREFERRED EMBODIMENTS OF THE INVENTION

The drug substance of the formulation according to the present invention is a 5-acetoxy or hydroxy-4,6-di-tert-butyl-2,3-dihydrobenzofuran derivative represented by the above general formula (1), which is known as described in WO94/08930, or probucol. BO-653 is preferable. As used herein, BO-653 refers to 4,6-di-tert-butyl-2,2-di-n-pentyl-5-hydroxy-2,3-dihydrobenzofuran.

In the definition of the compounds of general formula (1), the examples of the acyl group represented by R_1 include acetyl, formyl, propionyl, benzoyl and benzyloxycarbonyl groups.

The lower alkyl group represented by R_2 means a straight or branched alkyl group having 1 to 6 carbon atoms, the examples of which include, for example, methyl, ethyl, n-propyl, i-propyl, n-butyl, s-butyl and t-butyl groups.

The alkyl group represented by R_3 and R_4 means a straight or branched alkyl group having 1 to 20 carbon atoms, the examples of which include, for example, methyl, ethyl, n-propyl, i-propyl, n-butyl, s-butyl, t-butyl, pentyl, hexyl, heptyl, octyl, nonyl and decyl groups.

The alkenyl group represented by R_3 and R_4 means a straight or branched alkenyl group having 2 to 20 carbon atoms, the examples of which include, for example, vinyl, allyl, butenyl, pentenyl, geranyl and farnesyl groups.

The alkynyl group represented by R_3 and R_4 means a straight or branched alkynyl group having 2 to 20 carbon atoms, the examples of which include, for example, ethynyl,

propynyl and butynyl groups.

5 The aryl group represented by R_3 and R_4 means a monovalent aromatic hydrocarbon formed by removing one hydrogen atom from the aromatic hydrocarbon, the examples of which include, for example, phenyl, tolyl, xylyl, biphenyl, naphthyl, anthryl and phenanthryl groups. The ring carbon atoms of the aryl group may be substituted by one or more groups selected from halogen, lower alkyl, hydroxyl, alkoxy, amino, nitro or trifluoromethyl.

10 The examples of substituents on these alkyl, alkenyl, alkynyl and aryl groups represented by R_3 and R_4 include halogen, hydroxyl, amino, alkoxy and aryloxy.

The examples of the fat and oil according to the present invention include vegetable oils, medium chain fatty acid triglycerides (hereinafter referred to as MCTs), tricaprylin, oleic acid, linoleic acid, linolenic acid and lecithin. Vegetable oils are preferable.

Preferred vegetable oils are those containing an unsaturated fatty acid as a main component, the examples of which include, for example, soybean oil, safflower oil, sunflower oil, corn oil, cottonseed oil, sesame oil, rapeseed oil, peanut oil and olive oil, among which soybean oil is especially preferred. As used herein, the term "unsaturated fatty acid" means a fatty acid having one or more double bonds in its molecule, the examples of which include, for example, oleic acid, linoleic acid and linolenic acid. To be regarded as the main component, the unsaturated fatty acid should be present in the vegetable

oil in an amount of 50% or more of the total amount of the vegetable oil.

5 The amount of the fat and oil which may be present in the formulation of this invention is not limited so far as it enables the derivative or probucol to be formulated with the fat and oil. However, the fat and oil may be present in the formulation in an amount of 10-98% by weight based on the total weight of the filling solution.

As used herein, the term "soft capsule formulation" means a formulation in which a solution of a drug in a fat and oil is encapsulated in a capsule shell made of a gelatin-based material. The term "seamed soft capsule" , as used herein, means the soft capsule which is prepared by a rotary method. The rotary method is for producing a soft capsule and comprises the steps of stamping capsule shells out of a gelatin sheet which has been formed by gelling a solution containing gelatin, plasticizer and purified water, and then fusing the edges of the capsule shells each other with injecting filling solution therein to form a soft capsule. Hence, this method is also called "stamping method". The formed soft capsule has a seam due to the fusing step.

The seamed soft capsule shell of the present invention contains various types of gelatin and plasticizer. 25 The gelatins include those derived from animals such as cattle and swine. The examples of type of gelatin include an alkali-treated gelatin, an acid-treated gelatin, and a chemically modified gelatin. The gelatin may be used alone

or as a mixture of two or more.

The alkali-treated gelatins are those obtained by hydrolyzing a raw material of gelatin such as collagen or ossein with an alkaline material such as a lime solution
5 and then extracting the hydrolyzate, while the acid-treated gelatins are those obtained by hydrolyzing a collagen with an acidic material such as dilute hydrochloric acid or dilute sulfuric acid. The chemically modified gelatins include, but not limited to, those prepared by reacting an
10 amino group of a gelatin with succinic acid, phthalic acid, acetic acid or the like. The alkali-treated or acid-treated gelatins may be used as a starting material to prepare the chemically modified gelatins.

Suitable plasticizers include concentrated glycerin,
15 sorbitol, maltose, glucose, maltitose, sucrose, xylitol, mannitol, erythritol, polyethylene glycols having a molecular weight of 400-6000, etc.

The seamed soft capsule shell of the present invention preferably contains both gelatin and concentrated
20 glycerin.

When concentrated glycerin is used as a plasticizer, it is preferably combined with gelatin at a ratio of 15-50 parts by weight, more preferably 20-45 parts by weight to
100 parts by weight of the gelatin.

25 The seamed soft capsule shell of the present invention may contain further plasticizers such as sorbitol, maltose, glucose, maltitose, sucrose, xylitol, mannitol and erythritol. In the present invention, these are added to

suppress the deterioration, such as inteneration, stickiness and disintegration delay, of the shell.

Sorbitol is preferred among all. Preferably, sorbitol is present in the capsule shell at a ratio of 5-15 parts by weight, more preferably 7-10 parts by weight to 100 parts by weight of gelatin.

The seamed soft capsule shell preferably has a thickness of $28-40 \times 10^{-3}$ inch ($71-100 \times 10^{-2}$ mm), more preferably $30-35 \times 10^{-3}$ inch ($76-89 \times 10^{-2}$ mm) as formed.

10 The term "thickness as formed" is regarded as the thickness of a gelatin sheet out of which the shells are stamped. The term "thickness as formed" is hereinafter referred to as "thickness".

The examples of the formulation of the present invention include those in which a solution of BO-653 in a fat and oil, preferably a vegetable oil, more preferably soybean oil, containing unsaturated fatty acids as major constituent is filled into a seamed soft capsule, the shell of which preferably contains gelatin, concentrated glycerin and sorbitol.

15 20

The drug substance used for preparing the formulation of the present invention can be produced by known method. The compounds represented by general formula (1) including BO-653 can be produced by the method described for example in WO94/08930. Probuco1 can also be prepared by known method.

25

Fat and oils used for dissolving the drug substance therein, and the gelatins and plasticizers described above

which are main components of the seamed soft capsule shells are all commercially available.

The formulations of the present invention can be prepared by known method. For example, the present
5 formulations may be prepared via the steps of: preparing a filling solution, preparing capsule shells, forming a seamed soft capsule by encapsulating the filling solution with the shells, and drying the formed capsule.

The step of preparing the filling solution comprises
10 mixing a drug substance and a fat and oil. The step of preparing capsule shells comprises mixing a gelatin, a plasticizer and the like.

The step of forming a seamed soft capsule is carried out by the rotary process. The rotary process comprises:
15 (1) passing a continuous gelatin sheet between two rotating rollers opposed to each other and stamping capsule shells out of the gelatin sheet by the action of rollers; and (2) at the same time as the stamping, injecting the filling solution into a pair of the capsule shells and then fusing
20 the edges of the shells by heat action to form a seamed soft capsule.

The drying step comprises drying the seamed soft capsule at about 25°C in the air controlled at a relative air humidity of about 5-10%.

25 The invention will be illustrated in more detail with reference to the following Examples, but the present invention is not deemed to be limited thereto. Test results demonstrating the utility of the present invention

will also be shown.

EXAMPLES

Example 1: Preparation of seamed soft capsule formulation 5 containing BO-653

Seamed soft capsule formulations 1-4 below were prepared by the method described above.

Formulation 1

10 Filling solution: 10% w/w solution of BO-653 in soybean oil.

Capsule shells: alkali-treated gelatin/concentrated glycerin = 100/30 w/w.

Formulation 2

15 Filling solution: 10% w/w solution of BO-653 in MCT.
Capsule shells: alkali-treated gelatin/concentrated glycerin = 100/30 w/w.

Formulation 3

20 Filling solution: 50% w/w solution of BO-653 in soybean oil.
Capsule shells: as shown in Table 1.

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Table 1: Capsule shells

Capsule No.	Thickness		Content of concentrated glycerin (per 100 parts by weight of gelatin)
	$\times 10^{-3}$ inch	$\times 10^{-2}$ mm	
1	26	66	30.0
2	30	76	30.0
3	34	86	30.0
4	26	66	44.2
5	30	76	44.2
6	34	86	44.2
7	26	66	56.4
8	30	76	56.4
9	34	86	56.4

Dosage form: 5 oval

Gelatin: alkali-treated gelatin

Amount of solution filled: 250 mg/capsule.

5 Formulation 4

Filling solution: 4, 40 and 80% w/w solutions of BO-653 in soybean oil.

Capsule shells: as shown in Table 2.

Table 2

Content of concentrated glycerin	20 (per 100 parts by weight of gelatin)
Content of sorbitol	7 (per 100 parts by weight of gelatin)
Thickness	30×10^{-3} inch (76×10^{-2} mm)

10 Gelatin: alkali-treated gelatin.

Test example 1: Stability of BO-653 in various fat and oils

5 The influence of oxygen on the stability of BO-653 in various fat and oils was tested. BO-653 was dissolved in soybean oil, olive oil, MCT, tricaprylin, and linoleic acid separately to give a concentration of 50% by weight. 70 mg of each of the solutions was placed into individual 5 cc glass bottles. Separately, 35 mg of BO-653 was directly placed into a 5 cc glass bottle. The samples were prepared in duplicate. One set of samples were subjected to an argon replacement of the headspace of the bottles and then tightly sealed, while the other set of samples were sealed without argon replacement. Accelerated test at 80°C was conducted using all of the samples. The residual ratio of BO-653 in each bottle after 17 hours was determined by HPLC under the conditions shown in Table 3.

Table 3: HPLC analysis

Column	DEVELOSIL ODS-HG-5, 250 mm x 4.6 mm I.D. (Nomura Chemical)
Mobile phase	Acetonitrile/2-propanol/water (800:120:80)
Flow rate	1 mL/min
Column temperature	Room temperature
Detection wavelength	300 nm

The test results are shown in Fig. 1.

20 For the samples, including one containing only BO-653, with the argon replacement (i.e., in the absence of oxygen), little loss of BO-653 was observed indicating that BO-653

is stable in any fat and oils used. For the samples without the argon replacement (i.e, in the presence of oxygen), the stability of BO-653 varied depending on the fat and oils used. Relatively high stability was observed in the case where soybean oil was used.

Test example 2: Influence of administration solvents on the capability of being absorbed by body of BO-653

In order to determine the influence of administration solvents on the capability of being absorbed by body of BO-653, the following tests were conducted using the samples prepared with different solvents shown in Table 4.

Table 4: Samples

Sample 1	BO-653 solution in soybean oil having a concentration of 25 mg/mL
Sample 2	BO-653 solution in MCT having a concentration of 25 mg/mL
Sample 3	BO-653 suspension in 3% aqueous solution of gum arabic having a BO-653 concentration of 25 mg/mL

8-week old SD male rats (270-300 g) were divided into several groups of four. Each of the samples was orally administered via a stomach catheter to the animals under fasting so that a dose of BO-653 of 50 mg/kg was given. After a certain period, blood was collected from the tail vein, and plasma was separated and then measured for plasma BO-653 levels by HPLC under conditions shown in Table 5.

Table 5: HPLC analysis

Column	TSK-gel ODS-80Ts, 2.0 mm x 150 mm (TOSOH)
Mobile phase	Methanol/water (95:5)
Flow rate	200 μ L/min
Column temperature	Room temperature
Detection wavelength	300 nm

The test results are shown in Table 6 and Fig. 2.

Table 6: Pharmacokinetic parameters (mean \pm SD, n = 4)

Sample	C_{max} in μ g/mL	AUC_{0-48h} in μ g.h/mL
Sample 1	6.1 \pm 3.5	44.6 \pm 15.7
Sample 2	2.1 \pm 0.5	10.6 \pm 4.3
Sample 3	2.2 \pm 0.3	23.2 \pm 7.3

5

In Table 6, C_{max} represents the peak plasma level and AUC_{0-48h} represents the area under the plasma level-time curves shown in Fig. 2 from 0 to 48 hours after administration. In Fig. 2, "soybean oil", "MCT" and "3% gum arabic" are for samples 1, 2 and 3, respectively. Sample 1 in which soybean oil was used as solvent showed the highest plasma level shift and the AUC value for Sample 1 was about 4 times and about twice as high as compared with those for samples 2 and 3, respectively.

15

Test example 3: Stability of seamed soft capsule formulation of BO-653 (1)

Formulations 1 and 2 obtained in Example 1 were

tested for their stability under the conditions below:

Condition 1:

Formulation 1 was left at 40°C in a 100% oxygen atmosphere. The content of BO-653 after 1 month was determined by HPLC under the conditions shown in Table 3.

Condition 2:

Formulation 1 was left at 50°C in a 100% oxygen atmosphere. The content of BO-653 after 2 weeks was determined by HPLC under the conditions shown in Table 3.

Condition 3:

Formulation 2 was left at 50°C in a 100% oxygen atmosphere. The content of BO-653 after 3 weeks was determined by HPLC under the conditions shown in Table 3.

As a control, BO-653 was dissolved in soybean oil to give a concentration of 10% by weight and the solution was left at 40°C in a 100% oxygen atmosphere in the form that the solution was not filled into a soft capsule. The residual ratio of BO-653 after 1 month was determined by HPLC under the conditions shown in Table 3. The test results are shown in Table 7.

Table 7: Residual ratio (%) of BO-653 after various accelerated tests

Condition 1	100.7
Control	< 0.2
Condition 2	99.6
Condition 3	92.8

As apparent from comparison between condition 1 and control, BO-653 shows very high stability when it is filled into a soft capsule. Comparison between condition 2 and condition 3 shows that the stability of BO-653 in a soft capsule is influenced by the fat and oil used and that the stability is especially high when BO-653 is dissolved in soybean oil.

Test example 4: Stability of seamed soft capsule
10 formulations of BO-653 (2)

Formulation 3 obtained in Example 1 was stored at 50°C in a 100% oxygen atmosphere. The residual ratio of BO-653 after 1 month was determined by HPLC under the conditions shown in Table 3. The test results are shown in
15 Table 8.

Table 8: Content (%) of BO-653 after left at 50°C in a 100% oxygen atmosphere for 1 month

Glycerin content	Thickness		
	26 x 10 ⁻³ inch (66 x 10 ⁻² mm)	30 x 10 ⁻³ inch (76 x 10 ⁻² mm)	34 x 10 ⁻³ inch (86 x 10 ⁻² mm)
30.0	99.4	99.9	99.7
44.2	98.8	100.0	100.8
56.4	-*1	97.0	98.8

*1 Not evaluated because the capsule shell melted.

BO-653 shows especially high stability when
20 encapsulated in a soft capsule shell having a thickness of 30-34 x 10⁻³ inch (76-86 x 10⁻² mm) and containing 30.0-44.2 parts by weight of concentrated glycerin per 100 parts by

weight of gelatin. Especially preferred thickness is 30-34 x 10⁻³ inch (76-86 x 10⁻² mm) because it allows the shell to have an excellent oxygen barrier property and a good disintegrating property. In order to improve the oxygen barrier property, the shell preferably contains 30 parts by weight of concentrated glycerin per 100 parts by weight of gelatin. Seamed soft capsule formulation 3 is considered to show an excellent stability and a good disintegrating property.

Test example 5: Stability of seamed soft capsule formulations of BO-653 (3)

Formulation 4 obtained in Example 1 was stored under various conditions shown in Table 9. The residual ratio of BO-653 after the storage was determined by HPLC under the conditions shown in Table 3. The test results are shown in Table 9.

Table 9: Residual ratio (%) of BO-653 after various accelerated tests

Accelerated test conditions	BO-653 content in filling solution		
	4% by weight	40% by weight	80% by weight
40°C, RH75% - 1M	99.0	100.1	99.8
50°C, 100% oxygen atmosphere-1M	100.0	100.7	101.0
Irradiation with 1,200,000 lux.hrs white light	100.0	100.0	99.5
Irradiation with 1,200,000 lux.hrs white light followed by 400 W.hrs/m ² UV rays	98.0	99.7	99.8

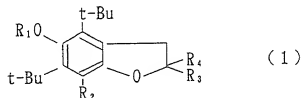
These formulations show an excellent stability under any storage conditions or at any BO-653 contents. The shells were neither intenerated nor sticky. It can be concluded from these results that Formulation 4, the shell of which contains sorbitol as a further component and has a lower glycerin content than that of Formulation 3, is more preferred seamed soft capsule formulation than any other formulations because it not only exhibits an excellent stability but also suppresses inteneration, stickiness, and disintegration delay of the shell.

INDUSTRIAL APPLICABILITY

The present invention provides BO-653 formulation having an excellent stability and a capability of being enterally absorbed.

CLAIMS

1. A seamed soft capsule formulation in which a solution of a dihydrobenzofuran derivative represented by general formula (1):



wherein R_1 represents a hydrogen atom or an acyl group; R_2 represents a hydrogen atom or a lower alkyl group; and R_3 and R_4 , which may be identical or different, represent a hydrogen atom or a substituted or unsubstituted alkyl, alkenyl, alkynyl or aryl group, or probucol in a fat and oil is filled into a seamed soft capsule.

2. The formulation of Claim 1 wherein said compound represented by general formula (1) is 4,6-di-tert-butyl-2,2-di-n-pentyl-5-hydroxy-2,3-dihydrobenzofuran.

3. The formulation of Claim 1 or 2 wherein said fat and oil is a vegetable oil.

4. The formulation of Claim 3 wherein said vegetable oil contains an unsaturated fatty acid as a major constituent.

5. The formulation of Claim 4 wherein said vegetable oil is soybean oil.

6. The formulation of Claim 1 or 2 wherein shell of said soft capsule contains gelatin and concentrated glycerin.

7. The formulation of Claim 1 or 2 wherein shell of

said soft capsule contains sorbitol.

8. The formulation of Claim 6 wherein said shell of said soft capsule further contains sorbitol.

9. The formulation of Claim 8 wherein said soft capsule contains 15-50 parts by weight of concentrated glycerin per 100 parts by weight of gelatin.

10. The formulation of Claim 9 wherein said soft capsule contains 20-45 parts by weight of concentrated glycerin per 100 parts by weight of gelatin.

11. The formulation of Claim 8 wherein said shell of said soft capsule contains 5-15 parts by weight of sorbitol per 100 parts by weight of gelatin.

12. The formulation of Claim 11 wherein said sorbitol is present in said shell at a ratio of 7-10 parts by weight of sorbitol to 100 parts by weight of gelatin.

13. The formulation of Claim 1 or 2 wherein a thickness as formed of shell of said soft capsule is from 28×10^{-3} inch (71×10^{-2} mm) to 40×10^{-3} inch (100×10^{-2} mm).

14. The formulation of Claim 13 wherein said thickness is from 30×10^{-3} inch (76×10^{-2} mm) to 35×10^{-3} in (89×10^{-2} mm).

15. A seamed soft capsule formulation containing 4,6-di-tert-butyl-2,2-di-n-pentyl-5-hydroxy-2,3-dihydrobenzofuran.

16. A method for treating an arteriosclerosis comprising administering said seamed soft capsule formulation of Claim 1 or 2 to a patient suffering from

said arteriosclerosis.

17. A method for treating an arteriosclerosis comprising administering said seamed soft capsule formulation of Claim 15 to a patient suffering from said arteriosclerosis.

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ABSTRACT

A seamed soft capsule formulation in which a solution
of BO-653, 4,6-di-tert-butyl-2,2-di-n-pentyl-5-hydroxy-2,3-
5 dihydrobenzofuran, in a fat and oil is filled into a seamed
soft capsule is provided. This BO-653 formulation confers
an excellent stability and a capability of being absorbed
by body to BO-653.

09914066.082301

Fig. 1

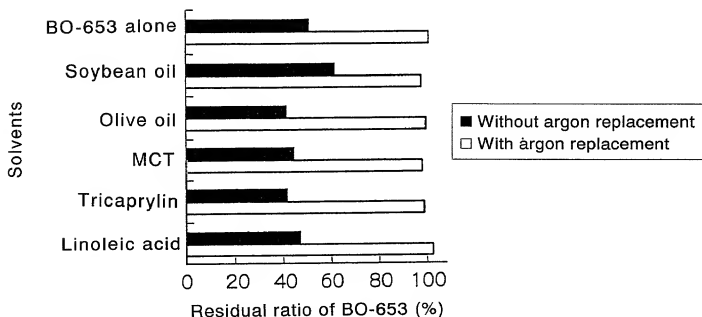
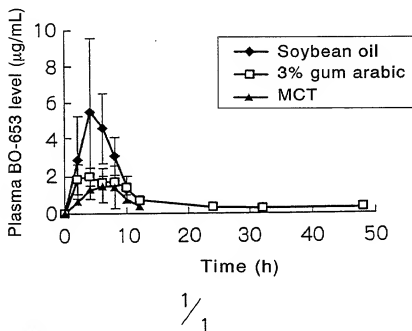


Fig. 2

(mean \pm SD, n = 4)1/
1

Combined Declaration for Patent Application and Power of Attorney

As a below-named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name; and that I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

SEAMED SOFT CAPSULE FORMULATION CONTAINING DIHYDROBENZOFURAN DERIVATIVE

the specification of which (check one)

- ☐ is attached hereto;
☐ was filed in the United States under 35 U.S.C. §111 on _____, as
 U.S. Appl. No. _____*; or
☒ was/will be filed in the U.S. under 35 U.S.C. §371 by entry into the U.S. national stage of an international (PCT) application, PCT/JP00/00862 filed Feb. 16, 2000, entry requested on _____*; national stage application received U.S. Appl. No. _____*; §371/§102(e) date _____* (* if known)

and was amended on _____ (if applicable).

(include dates of amendments under PCT Art. 19 and 34 if PCT)

I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above; and I acknowledge the duty to disclose to the Patent and Trademark Office (PTO) all information known by me to be material to patentability as defined in 37 C.F.R. §1.56.

I hereby claim foreign priority benefits under 35 U.S.C. §§ 119 and 365 of any prior foreign application(s) for patent or inventor's certificate, or prior PCT application(s) designating a country other than the U.S., listed below with the "Yes" box checked and have also identified below any such application having a filing date before that of the application on which priority is claimed:

<u>45208/1999</u>	<u>Japan</u>	<u>23/2/99</u>	<input checked="" type="checkbox"/> <input type="checkbox"/>
(Number)	(Country)	(Day Month Year Filed)	YES NO
_____	_____	_____	<input type="checkbox"/> <input type="checkbox"/>
(Number)	(Country)	(Day Month Year Filed)	YES NO

I hereby claim the benefit under 35 U.S.C. §120 of any prior U.S. non-provisional application(s) or prior PCT application(s) designating the U.S. listed below, or under §119(c) of any prior U.S. provisional applications listed below, and, insofar as the subject matter of each of the claims of this application is not disclosed in such U.S. or PCT application in the manner provided by the first paragraph of 35 U.S.C. §112, I acknowledge the duty to disclose to the PTO all information as defined in 37 C.F.R. §1.56(a) which occurred between the filing date of the prior application and the national filing date of this application:

_____ (Application No.)	_____ (Day Month Year Filed)	_____ (Status: patented, pending, abandoned)
_____ (Application No.)	_____ (Day Month Year Filed)	_____ (Status: patented, pending, abandoned)
_____ (Application No.)	_____ (Day Month Year Filed)	_____ (Status: patented, pending, abandoned)

As a named inventor, I hereby appoint the following registered practitioners to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

All of the practitioners associated with Customer Number 001444

Direct all correspondence to the address associated with Customer Number 001444; i.e.,
BROWDY AND NEIMARK, P.L.L.C.
 624 Ninth Street, N.W.
 Washington, D.C. 20001-5303
 (202) 628-5197

The undersigned hereby authorizes the U.S. Attorneys or Agents appointed herein to accept and follow instructions from YUASA AND HARA as to any action to be taken in the U.S. Patent and Trademark Office regarding this application without direct communication between the U.S. Attorneys or Agents and the undersigned. In the event of a change of the persons from whom instructions may be taken, the U.S. Attorneys or Agents appointed herein will be so notified by the undersigned.

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